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Sheet 1 of 2

Complete if Known	
Application Number	10/021,955
Filing Date	December 13, 2001
First Named Inventor	James R. Lupski
Art Unit	1645
Examiner Name	Not Yet Assigned
Attorney Docket Number	HO-P02086US1

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Document Number: Number-Kind Code[2] *(if known)*	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear

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NON PATENT LITERATURE DOCUMENTS

Examiner Initials*	Cite No.[1]	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T[2]
gpc	CA	Guilbot, Angele, et al.; A mutation in periaxin is responsible for CMT4F, an autosomal recessive form of Charcot-Marie-Tooth disease, Human Molecular Genetics, (2001) Vol. 10., No. 4, pp. 415-421.	
	CB	Delague, Valerie, et al.; Mapping of a new locus for autosomal recessive demyelinating charcot-marie-tooth disease to 19q13.1-13.3 in a large consanguineous lebanese family: exclusion of MAG as a candidate gene, Am. J. Hum. Genet. (2000) 67:236-243.	
	CC	Williams, Anna C., et al.; The function of the periaxin gene during nerve repair in a model of CMT4F*; J. Anat. (2002) 200, pp. 323-330.	
	CD	Boerkoel, Cornelius F., et al.; Charcot-marie-tooth disease and related neuropathies: mutation distribution and genotype-phenotype correlation, Annals of Neurology (2002) 51:190-201.	
	CE	Zhou, L., Griffin JW; Demyelinating neuropathies, Curr. Opin. Neurol. (June 2003), 16:(3):307-313.p	
	CD	Martin, J.J., et al.; Neuropathology of some hereditary conditions affecting central and peripheral nervous system; Acta Neurol. Belg. (2002), 102, pp. 30-35.	

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25366913.1 Examiner	Prabha Chandran	Date Considered	4/29/04